

# FISHing Cuts the Angst in Amniocentesis

*A new technique produces results in days instead of weeks; soon, amniocentesis itself may be superseded*

ONE OF THE MOST WRENCHING TIMES A couple can face is the week—or two or three—that they must wait for the results of amniocentesis, which will tell them whether the baby they are expecting will have a serious birth defect such as Down's syndrome. Now a research team at Integrated Genetics, a biotechnology company in Framingham, Massachusetts, has come up with a new technique that can slash the waiting time from 3 weeks to less than 2 days. And two other groups are working on an alternative that may eventually make both amniocentesis and chorionic villus sampling (CVS), another prenatal test, obsolete.

Most of the delay in amniocentesis comes from the time it takes to grow the fetal cells in culture so that a sufficient number of them will be in metaphase—the time during the cell cycle when the chromosomes can be distinguished from one another visually. In the standard approach, cytogeneticists then examine the distinctive bands on each chromosome to look for the telltale signs of chromosome breakage, duplication, or loss.

Integrated Genetics' new approach, described last week at the International Congress of Human Genetics in Washington, D.C., by geneticist Katherine Klinger, analyzes the nuclear DNA directly when the cells are in another phase, interphase, and the chromosomes look "like a big bowl of spaghetti." To do so, Klinger's group developed sets of DNA probes that home in on regions of five chromosomes—21, 18, 13, X, and Y—which together account for 90% to 95% of all birth defects related to chromosomal abnormalities. The probes are then tagged with fluorescent dyes that glow different colors when examined under ultraviolet light. In a normal cell, for instance, they would expect to see that two brightly colored dots, indicating two copies of chromosome 21, are present. If they see three, the fetus has Down's syndrome.

In a double-blind clinical trial, the Integrated Genetics group analyzed 500 fetal samples and compared their results with those obtained from standard cytogenetic analysis. Their method, called FISH (fluorescent in situ hybridization), correctly identified all 21 abnormalities in the samples, without any false positives or negatives, reported Klinger.

Klinger views the test as "an enhancement to, not a replacement for" the standard methods of chromosome analysis. In fact, Integrated Genetics won't offer the test unless the couple agrees to have the full analysis done as well. "This gives the couple the best of both worlds—rapid determination of the major defects, followed up by complete karyotyping [analysis] to ensure that the less common chromosome disorders are also detected," Klinger says.

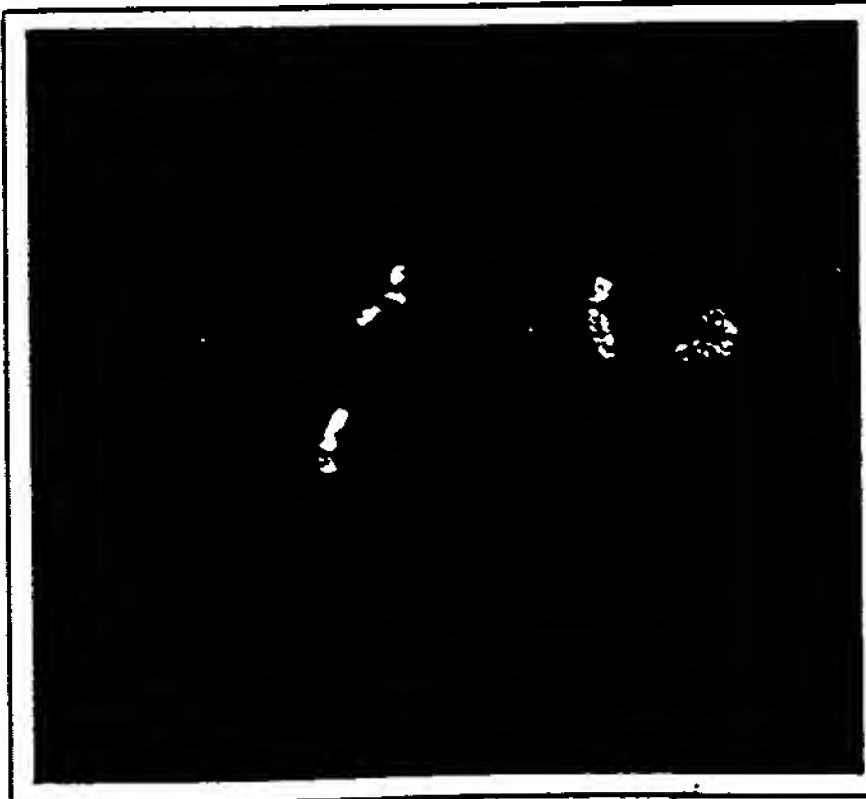
The next step is to develop probe sets that will reveal the rarer abnormalities, such as the tiny chromosome deletions involved in Prader-Willi syndrome, that are difficult to detect with existing techniques. But the ultimate power of this approach, Klinger says, will be in its application in noninvasive alternatives to both amniocentesis and CVS. Klinger is in a good position to know, as she is collaborating with two of the groups working on such techniques, one led by Diana Bianchi of Children's Hospital in Boston, and the other by Stephen Wachtel and Sherman Elias of the University of Tennessee, Memphis. Both groups described their progress in Washington.

A major problem with both amniocentesis and CVS—aside from the waiting time—is the risk of miscarriage, estimated at 1% to 2% for CVS and at 0.5% for amniocentesis. Both the Boston and Memphis groups now have evidence that it may be possible to

## Paint Kit for Cancer Diagnosis

Biophysicists Joe Gray and Dan Pinkel have spent much of the past decade perfecting a way to "paint" entire chromosomes so that abnormalities associated with cancer stand out. Last week, at a press conference on the opening day of the International Congress of Human Genetics in Washington, D.C., they unveiled the fruits of their labor: a chromosome painting kit that will be marketed for research use later this month by Imagenetics, an Illinois-based medical diagnostics company. Imagenetics has dubbed the technology Whole Chromosome Paints.

Gray and Pinkel, both of whom moved to the University of California at San Francisco in July, developed their painting technique while at the Lawrence Livermore National Laboratory. It is based on fluorescent in situ hybridization (see story), in which fluorescently labeled probes are used to stain, or paint, entire chromosomes. The "paints" consist of anywhere from 10,000 to 100,000 fragments of DNA for each of the 24 chromosomes. When these fragments are tagged with a fluorescent dye and hybridized to the chromosome, the entire chromosome lights up, making any abnormalities readily apparent to even an untrained eye. If, for example, a piece of chromosome 9 has been exchanged with a portion of chromosome 22, as often happens in chronic myelogenous leukemia, the abnormality would be clearly visible as, say, a stretch of green in a red chromosome, and vice versa. To David Galas, director of health and environmental research at the Department of Energy, which supported some of this work, the biggest appeal of this approach is its potential for automation, which is "almost impossible" in conventional chromosome analysis.



**Paints.** The new kit reveals that chromosome 12 (green) and chromosome 7 (red) have exchanged pieces.

In research supported by Imagenetics, chromosome paints are now being tested in several laboratories. At the Mayo Clinic, for instance, Robert Jenkins is using them to analyze cells for the presence of three copies of chromosome 8, a common abnormality in leukemia. At Livermore, two groups are using them to look for chromosome damage resulting from exposure to chemicals and radiation. And in the future, says Gray, this technique will be extended to scan for more subtle abnormalities in specific oncogenes or tumor suppressor genes that may underlie the development of cancer. Indeed, such work is already getting under way. ■ L.R.

sidestep that risk entirely by obtaining fetal cells from the mother's blood.

The first step was to confirm that fetal cells are actually present in the mother's blood—though in exceedingly low amounts—as several groups had suggested. The researchers did that by using polymerase chain reaction (PCR) techniques to detect and amplify a specific sequence of DNA from the Y chromosome, which presumably would come only from a male fetus. They were able to predict the sex of the child with 65% accuracy—"not an exciting number," as Wachtel concedes, but enough to convince them that fetal cells were indeed there.

Their goal was to find a way to enrich the concentration of those cells to get enough for prenatal diagnosis. For that the two groups used a flow cytometer, an instrument that uses laser light to separate cells according to different characteristics—in this case, on the basis of cell size, granularity, and the presence of two cell surface markers characteristic of fetal cells. The researchers then performed the same PCR analysis to see how well the enrichment worked: This time they were able to predict sex with 94% accuracy. Flow cytometry clearly works, says Wachtel, enriching the concentration of fetal cells from an estimated 1 in 20 million maternal cells to about 1 in 100.

Both groups then sent off their sorted fetal cells, fixed on glass slides, to Klinger to see whether they could actually be used for prenatal diagnosis with the new FISH technique. One slide from Wachtel and Elias' group contained cells from a woman who they knew, from CVS, was carrying a fetus with Klinefelter syndrome, a rare disorder in which males have two copies of the X chromosome. "When Kathy called and said we see an XXY male, we started to get excited," Wachtel recalls. Integrated Genetics also accurately detected a fetus with Down's syndrome from both groups.

But the "big splash," as Wachtel calls it, came when Wachtel's group sent a sorted sample from a 42-year-old woman who had not had CVS or amniocentesis. Integrated Genetics found the fetus carried three copies of chromosome 18, a syndrome that leads to severe mental retardation. That was the first time a prenatal disorder was actually diagnosed, rather than just confirmed, through this new approach, Wachtel says. He cautions, however, that "this is the first shot. It is certainly not ready for routine use yet." Indeed, his group has analyzed samples from just 41 women—far too few to determine either the specificity or sensitivity of the procedure. Klinger expects clinical trials to be under way within 2 years. "That is clearly where all of us would like prenatal diagnosis to go." ■ LESLIE ROBERTS

## Brave New (RNA) World

**Cambridge**—The earliest days of life, many researchers now say, may have been played out in an "RNA world," in which one large molecule, RNA, carried out all the processes of life. But as acceptance of that idea spreads, its proponents are facing a new problem: How did the RNA world give way to the more complex biochemistry of life as we know it? That conundrum emerged as a key theme of Biological Functions of RNA, a symposium sponsored last week by the Whitehead Institute for Biomedical Research that was chock full of provocative notions.

The starting point for these notions—the theory of an RNA world—grew out of researchers' frustration at the seemingly neat division of labor in the realm of biomolecules. Protein enzymes do the heavy lifting, catalyzing the chemical reactions needed to sustain and reproduce life. DNA and RNA have it easier, serving mainly as a medium for the genetic information that guides all that heavy lifting. Life requires both functions, which seemed to imply that molecules filling both roles must somehow—defying all probability—have appeared at the same time in early evolution. But how?

That puzzle seemed on the way to being solved several years ago, when researchers found that RNA can play both roles: Besides storing genetic information, it can act as an enzyme. With that discovery, RNA began looking like a good candidate for being the primordial living molecule. But how would a swarm of specialized RNA enzymes have given way to a breed of proteins subsuming RNA's biochemical functions? At the Whitehead symposium, researchers presented startling evidence that RNA enzymes might well have fostered the transition by filling key niches in the world of proteins.

Molecular geneticist Alan Lambowitz of Ohio State University argued, for example, that one particular molecule that is a linchpin in the process of making new proteins may have evolved from an RNA enzyme. The evidence comes from a comparison between a present-day RNA enzyme found in the yeast *Neurospora crassa* and transfer RNA (tRNA), the linchpin molecule in question. tRNA is a shuttle that carries amino acids—the building blocks of proteins—to protein factories called ribosomes. By chance, Lambowitz and his colleagues found that a protein that binds to present-day tRNA molecules and attaches amino acids to them can also bind to part of the *Neurospora* RNA.

Lambowitz' interpretation of the result: Both RNA molecules must have the same three-dimensional shape, even though their sequences are very different. And that suggests to Lambowitz that some precursor of the RNA enzyme evolved into tRNA, and was thus recruited for protein synthesis.

Thomas Cech of the University of Colorado, a codiscoverer of catalytic RNA, raised the possibility of an even more active role for RNA enzymes in the primordial protein world: in the very synthesis of proteins. Doing so would require RNA to catalyze the formation of the specific bonds, called peptide bonds, that join amino acids in a protein. That, in turn, would imply that RNA could interact chemically with the carbon atoms in amino acids. But so far RNA's ability to make and break bonds has seemed to be confined to the bonds joining phosphorus and oxygen in RNA itself.

Now work by Joe Piccirilli in Cech's laboratory has shown that an RNA enzyme can break a bond between an amino acid and a nucleic acid, which requires an interaction between the RNA enzyme and the carbon in the amino acid. And if RNA can break such bonds, says Cech, maybe it can make them as well, which would open the possibility of RNA-catalyzed protein synthesis at some point in evolution.

Indeed, RNA-driven protein synthesis may be going on even now, in some present-day ribosomes, according to biochemist Harry Noller of the University of California, Santa Cruz. Within the complex of proteins and RNA that makes up a ribosome, it has been generally assumed that the protein enzymes actually do the catalytic work of forming bonds between amino acids, while the RNA serves as a structural rack for those proteins. But Noller's results suggest the ribosomal RNA may turn out to have the glamour role after all. Remarkably, Noller found that even when he teased away almost all of the protein from the ribosome of a bacterium, the ribosome was still highly effective at assembling amino acids. Noller stresses that he won't know for sure that RNA, and not protein, is actually catalyzing protein synthesis until he can demonstrate that ribosomal RNA completely denuded of proteins is still capable of catalyzing the reaction. But if he succeeds, Noller may have shown that, in one important respect, we are living in an RNA world even today.

■ MICHELLE HOFFMAN